



(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
14.11.2001 Bulletin 2001/46

(51) Int Cl.7: **H01L 21/314**

(21) Application number: **01102401.5**

(22) Date of filing: **02.02.2001**

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: **17.02.2000 US 183507 P**
08.06.2000 US 590322

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(54) **Method of depositing an amorphous carbon layer**

(57) A method of forming an integrated circuit using an amorphous carbon film. The amorphous carbon film is formed by thermally decomposing a gas mixture comprising a hydrocarbon compound and an inert gas. The amorphous carbon film is compatible with integrated circuit fabrication processes. In one integrated circuit fab-

rication process, the amorphous carbon film is used as a hardmask. In another integrated circuit fabrication process, the amorphous carbon film is an anti-reflective coating (ARC) for deep ultraviolet (DUV) lithography. In yet another integrated circuit fabrication process, a multi-layer amorphous carbon anti-reflective coating is used for DUV lithography.

Description

[0001] The present invention relates to an amorphous carbon film, its use in integrated circuit fabrication, and a method for depositing the film.

[0002] Integrated circuits have evolved into complex devices that can include millions of transistors, capacitors and resistors on a single chip. The evolution of chip designs continually requires faster circuitry and greater circuit density. The demands for faster circuits with greater circuit densities impose corresponding demands on the materials used to fabricate such integrated circuits. In particular, as the dimensions of integrated circuit components are reduced (e. g., sub-micron dimensions), it has been necessary to use low resistivity conductive materials (e. g., copper) as well as low dielectric constant insulating materials (dielectric constant less than about 4.5) to improve the electrical performance of such components.

[0003] The demands for greater integrated circuit densities also impose demands on the process sequences used for integrated circuit manufacture. For example, in process sequences using conventional lithographic techniques, a layer of energy sensitive resist is formed over a stack of material layers on a substrate. An image of a pattern is introduced into the energy sensitive resist layer. Thereafter, the pattern introduced into the energy sensitive resist layer is transferred into one or more layers of the material stack formed on the substrate using the layer of energy sensitive resist as a mask. The pattern introduced into the energy sensitive resist can be transferred into one or more layers of the material stack using a chemical etchant. The chemical etchant is designed to have a greater etch selectivity for the material layers of the stack than for the energy sensitive resist. That is, the chemical etchant etches the one or more layers of the material stack at a much faster rate than it etches the energy sensitive resist. The faster etch rate for the one or more material layers of the stack typically prevents the energy sensitive resist material from being consumed prior to completion of the pattern transfer.

[0004] However, demands for greater circuit densities on integrated circuits have necessitated smaller pattern dimensions (e. g., sub-micron dimensions). As the pattern dimensions are reduced, the thickness of the energy sensitive resist must correspondingly be reduced in order to control pattern resolution. Such thinner resist layers (less than about 6000 Å) can be insufficient to mask underlying material layers during a pattern transfer step using a chemical etchant.

[0005] An intermediate oxide layer (e.g., silicon dioxide, silicon nitride), called a hardmask, is often used between the energy sensitive resist layer and the underlying material layers to facilitate pattern transfer into the underlying material layers. However, some material structures (e.g., damascene) include silicon dioxide and/or silicon nitride layers. Such material structures

cannot be patterned using a silicon dioxide or silicon nitride hardmask as an etch mask.

[0006] Resist patterning problems are further compounded when lithographic imaging tools having deep ultraviolet (DUV) imaging wavelengths (e. g., less than about 250 nanometers (nm)) are used to generate the resist patterns. The DUV imaging wavelengths improve resist pattern resolution because diffraction effects are reduced at these shorter wavelengths. However, the increased reflective nature of many underlying materials (e. g., polysilicon and metal silicides) at such DUV wavelengths, can degrade the resulting resist patterns.

[0007] One technique proposed to minimize reflections from an underlying material layer uses an anti-reflective coating (ARC). The ARC is formed over the reflective material layer prior to resist patterning. The ARC suppresses the reflections off the underlying material layer during resist imaging, providing accurate pattern replication in the layer of energy sensitive resist.

[0008] A number of ARC materials have been suggested for use in combination with energy sensitive resists. For example, U. S. Patent 5,626,967 issued May 6, 1997 to Pramanick et al. describes the use of titanium nitride anti-reflective coatings. However, titanium nitride is increasingly metallic as the exposure wavelength is reduced below 248 nm, meaning titanium nitride has high reflectivity for DUV radiation and is not an effective anti-reflective coating for DUV wavelengths.

[0009] U. S. Patent 5,710,067 issued January 20, 1998 to Foote et al. discloses the use of silicon oxynitride anti-reflective films. Silicon oxynitride films are difficult to remove, in that they leave residues behind that potentially interfere with subsequent integrated circuit fabrication steps.

[0010] Therefore, a need exists in the art for a material layer useful for integrated circuit fabrication, which has good etch selectivity with oxides. Particularly desirable would be a material layer that is also an ARC at DUV wavelengths and is easy to strip.

[0011] The present invention provides a method for forming an amorphous carbon layer for use in integrated circuit fabrication. The amorphous carbon layer is formed by thermally decomposing a gas mixture comprising a hydrocarbon compound and an inert gas. The gas mixture, which may optionally include an additive gas, is introduced into a process chamber where plasma enhanced thermal decomposition of the hydrocarbon compound in close proximity to a substrate surface, results in deposition of an amorphous carbon layer on the substrate surface.

[0012] An as-deposited amorphous carbon layer, deposited according to the process of the invention, has an adjustable carbon:hydrogen ratio that ranges from about 10 % hydrogen to about 60 % hydrogen. The amorphous carbon layer also has a light absorption coefficient, k , that can be varied between about 0.1 to about 1.0 at wavelengths below about 250 nm, making it suitable for use as an anti-reflective coating (ARC) at

DUV wavelengths.

[0013] The amorphous carbon layer is compatible with integrated circuit fabrication processes. In one integrated circuit fabrication process, the amorphous carbon layer is used as a hardmask. For such an embodiment, a preferred process sequence includes depositing an amorphous carbon layer on a substrate. After the amorphous carbon layer is deposited on the substrate, an intermediate layer is formed thereon. A pattern is defined in the intermediate layer and transferred into the amorphous carbon layer. Thereafter, the pattern is transferred into the substrate using the amorphous carbon layer as a hardmask. Additionally, the pattern defined in the amorphous carbon hardmask can be incorporated into the structure of the integrated circuit, such as for example in a damascene structure.

[0014] In another integrated circuit fabrication process, the amorphous carbon layer is used as a single layer anti-reflective coating for DUV lithography. For such an embodiment, a preferred process sequence includes forming the amorphous carbon layer on a substrate. The amorphous carbon layer has a refractive index (n) in the range of about 1.5 to 1.9 and an absorption coefficient (k) in the range of about 0.1 to about 1.0 at wavelengths less than about 250 nm. The refractive index (n) and absorption coefficient (k) for the amorphous carbon ARC are tunable, in that they can be varied in the desired range as a function of the temperature and composition of the gas mixture during layer formation. After the amorphous carbon layer is formed on the substrate, a layer of energy sensitive resist material is formed thereon. A pattern is defined in the energy sensitive resist at a wavelength less than about 250 nm. Thereafter, the pattern defined in the energy sensitive resist is transferred into the amorphous carbon layer. After the amorphous carbon layer is patterned, such pattern is optionally transferred into the substrate.

[0015] In still another integrated circuit fabrication process, a multi-layer amorphous carbon anti-reflective coating is used for DUV lithography. For such an embodiment a preferred process sequence includes forming a first amorphous carbon layer on a substrate. The first amorphous carbon layer has an index of refraction in the range of about 1.5 to about 1.9 and an absorption coefficient (k) in the range of about 0.5 to about 1.0 at wavelengths less than about 250 nm. After the first amorphous carbon layer is formed on the substrate, a second amorphous carbon layer is formed thereon. The second amorphous carbon layer has an index of refraction of about 1.5 to about 1.9 and an absorption coefficient in the range of about 0.1 to about 0.5. The refractive index (n) and absorption coefficient (k) for the first and second amorphous carbon layers are tunable, in that they can be varied in the desired range as a function of the temperature and composition of the gas mixture during layer formation. A layer of energy sensitive resist material is formed on the second amorphous carbon layer. A pattern is defined in the energy sensitive resist lay-

er at a wavelength less than about 250 nm. The pattern defined in the energy sensitive resist material is thereafter transferred into the second amorphous carbon layer followed by the first amorphous carbon layer. After the first amorphous carbon layer is patterned, such pattern is optionally transferred into the substrate.

[0016] An aspect of the invention is a method of forming an amorphous carbon layer on a substrate comprising positioning a substrate in a deposition chamber; providing a gas mixture to the deposition chamber, wherein the gas mixture comprises one or more hydrocarbon compounds and an inert gas; and heating the gas mixture to thermally decompose the one or more hydrocarbon compounds in the gas mixture to form an amorphous carbon layer on the substrate.

[0017] A preferred aspect of the above invention is a method wherein the amorphous carbon layer has an index of refraction in the range of about 1.5 to about 1.9.

[0018] A preferred aspect of the above invention is a method wherein the amorphous carbon layer has an absorption coefficient in the range of about 0.1 to about 1.0, at wavelengths less than about 250 nm.

[0019] An aspect of the invention is a method of forming a device, comprising forming one or more amorphous carbon layers on a substrate; and defining a pattern in at least one region of the one or more amorphous carbon layers.

[0020] A preferred aspect of the above invention is a method further comprising transferring the pattern defined in the at least one region of the one or more amorphous carbon layers into the substrate using the one or more amorphous carbon layers as a mask.

[0021] A preferred aspect of the above invention is a method further comprising the step of removing the one or more amorphous carbon layers from the substrate.

[0022] A preferred aspect of the above invention is a method wherein the substrate has one or more material layers formed thereon.

[0023] A preferred aspect of the above invention is a method wherein the oxide is selected from the group consisting of silicon dioxide, silicon nitride, silicon oxynitride, silicon carbide, and amorphous silicon.

[0024] A preferred aspect of the above invention is a method wherein the one or more amorphous carbon layers are removed from the substrate using an ozone, oxygen, or NH_3 plasma.

[0025] A preferred aspect of the above invention is a method wherein the one or more amorphous carbon layers each have an index of refraction in the range of about 1.5 to about 1.9.

[0026] A preferred aspect of the above invention is a method wherein the one or more amorphous carbon layers are formed on the substrate by positioning the substrate in a deposition chamber; providing a gas mixture to the deposition chamber, wherein the gas mixture comprises one or more hydrocarbon compounds and an inert gas; and heating the gas mixture to thermally decompose the one or more hydrocarbon compounds in

the gas mixture to form the one or more amorphous carbon layers on the substrate.

[0027] A preferred aspect of the above invention is a method wherein the one or more amorphous carbon layers each have a thickness in the range of about 50 Å to about 1500 Å.

[0028] A preferred aspect of the above invention is a method wherein the patterned radiation has a wavelength less than about 250 nanometers (nm).

[0029] An aspect of the invention is a method of fabricating a damascene structure, comprising forming a dielectric layer on a substrate; forming an amorphous carbon layer on the dielectric layer; patterning the amorphous carbon layer to define contacts/vias there-through; transferring the pattern formed in the amorphous carbon layer through the dielectric layer to form contacts/vias therein; removing the amorphous carbon layer from the patterned dielectric layer; and filling the contacts/vias formed in the dielectric layer with a conductive material.

[0030] A preferred aspect of the above invention is a method wherein the dielectric layer has a dielectric constant less than about 4.5.

[0031] A preferred aspect of the above invention is a method wherein the conductive material filling the contacts/vias has a resistivity less than about 5 $\mu\Omega$ -cm.

[0032] A preferred aspect of the above invention is a method wherein the dielectric layer is selected from the group consisting of silicon dioxide and fluorosilicate glass.

[0033] A preferred aspect of the above invention is a method wherein the conductive material filling the contacts/vias is selected from the group consisting of copper, aluminum, tungsten, and combinations thereof.

[0034] A preferred aspect of the above invention is a method wherein the one or more hydrocarbon compounds is selected from the group consisting of propylene (C_3H_6), propyne (C_3H_4), propane (C_3H_8), butane (C_4H_{10}), butylene (C_4H_8), butadiene (C_4H_6), or acetylene (C_2H_2) and combinations thereof.

[0035] A preferred aspect of the above invention is a method wherein the amorphous carbon layer has a carbon:hydrogen ratio in the range of about 10 % hydrogen to about 60 % hydrogen.

[0036] A preferred aspect of the above invention is a method wherein the inert gas is selected from the group consisting of helium, argon and combinations thereof.

[0037] A preferred aspect of the above invention is a method wherein the gas mixture further comprises an additive gas.

[0038] A preferred aspect of the above invention is a method wherein the additive gas is selected from the group consisting of ammonia, nitrogen, hydrogen, and combinations thereof.

[0039] A preferred aspect of the above invention is a method wherein the substrate is heated to a temperature between about 100 °C and about 500 °C.

[0040] A preferred aspect of the above invention is a

method wherein the deposition chamber is maintained at a pressure between about 1 Torr to about 20 Torr.

[0041] A preferred aspect of the above invention is a method wherein the gas mixture is provided to the deposition chamber at a flow rate in a range of about 50 sccm to about 500 sccm.

[0042] A preferred aspect of the above invention is a method wherein the electric field applied to the gas mixture is a radio frequency (RF) power.

[0043] A preferred aspect of the above invention is a method wherein the amorphous carbon layer has a thickness in the range of about 200 Å to about 1000 Å.

[0044] An aspect of the invention is a computer storage medium containing a software routine that, when executed, causes a general purpose computer to control a deposition chamber according to any of the above methods.

[0045] Embodiments of the present invention can be readily understood by considering the following detailed description in conjunction with the accompanying drawings, in which:

FIG. 1 depicts a schematic illustration of an apparatus that can be used for the practice of this invention;

FIGS. 2a-2e depict schematic cross-sectional views of a substrate structure at different stages of integrated circuit fabrication incorporating an amorphous carbon layer as a hardmask;

FIGS. 3a-3e depict schematic cross-sectional views of a damascene structure at different stages of integrated circuit fabrication incorporating an amorphous carbon layer as a hardmask;

FIGS. 4a-4c depict schematic cross-sectional views of a substrate structure at different stages of integrated circuit fabrication incorporating an amorphous carbon layer as an anti-reflective coating (ARC); and

FIGS. 5a-5d depict schematic cross-sectional views of a substrate structure at different stages of integrated circuit fabrication incorporating a multi-layer amorphous carbon ARC structure.

[0046] The present invention provides a method of forming an integrated circuit using an amorphous carbon layer. The amorphous carbon layer is formed by thermally decomposing a gas mixture comprising a hydrocarbon compound and an inert gas. The gas mixture, which may optionally include an additive gas, is introduced into a process chamber where plasma enhanced thermal decomposition of the hydrocarbon compound in close proximity to a substrate surface, results in deposition of an amorphous carbon layer on the substrate surface. The amorphous carbon layer is compatible with integrated circuit fabrication processes, discussed below.

[0047] FIG. 1 is a schematic representation of a wafer processing system 10 that can be used to perform amor-

phous carbon layer deposition in accordance with the present invention. This apparatus typically comprises a process chamber 100, a gas panel 130, a control unit 110, along with other hardware components such as power supplies and vacuum pumps. Details of the system 10 used in the present invention are described in a commonly assigned U.S. patent application, entitled "High Temperature Chemical Vapor Deposition Chamber", Serial No. 09/211,998, filed on December 14, 1998, and is herein incorporated by reference. The salient features of this system 10 are briefly described below. Examples of system 10 include CENTURA® systems, PRECISION 5000® systems and PRODUCER™ systems commercially available from Applied Materials Inc., Santa Clara, California.

[0048] The process chamber 100 generally comprises a support pedestal 150, which is used to support a substrate such as a semiconductor wafer 190. This pedestal 150 can typically be moved in a vertical direction inside the chamber 100 using a displacement mechanism (not shown). Depending on the specific process, the wafer 190 can be heated to some desired temperature prior to processing. In the present invention, the wafer support pedestal 150 is heated by an embedded heater element 170. For example, the pedestal 150 may be resistively heated by applying an electric current from an AC supply 106 to the heater element 170. The wafer 190 is, in turn, heated by the pedestal 150. A temperature sensor 172, such as a thermocouple, is also embedded in the wafer support pedestal 150 to monitor the temperature of the pedestal 150 in a conventional manner. The measured temperature is used in a feedback loop to control the power supply 16 for the heating element 170 such that the wafer temperature can be maintained or controlled at a desired temperature which is suitable for the particular process application. The pedestal 150 is optionally heated using a plasma or by radiant heat (not shown).

[0049] A vacuum pump 102, is used to evacuate the process chamber 100 and to maintain the proper gas flows and pressure inside the chamber 100. A showerhead 120, through which process gases are introduced into the chamber 100, is located above the wafer support pedestal 150. The showerhead 120 is connected to a gas panel 130, which controls and supplies various gases used in different steps of the process sequence.

[0050] The showerhead 120 and wafer support pedestal 150 also form a pair of spaced apart electrodes. When an electric field is generated between these electrodes, the process gases introduced into the chamber 100 are ignited into a plasma. Typically, the electric field is generated by connecting the wafer support pedestal 150 to a source of radio frequency (RF) power (not shown) through a matching network (not shown). Alternatively, the RF power source and matching network may be coupled to the showerhead 120, or coupled to both the showerhead 120 and the wafer support pedestal 150.

[0051] Plasma enhanced chemical vapor deposition (PECVD) techniques promote excitation and/or dissociation of the reactant gases by the application of the electric field to the reaction zone near the substrate surface, creating a plasma of reactive species. The reactivity of the species in the plasma reduces the energy required for a chemical reaction to take place, in effect lowering the required temperature for such PECVD processes.

[0052] In the present embodiment, amorphous carbon layer deposition is accomplished by plasma enhanced thermal decomposition of a hydrocarbon compound such as propylene (C_3H_6). Propylene is introduced into the process chamber 100 under the control of gas panel 130. The hydrocarbon compound is introduced into the process chamber as a gas with a regulated flow.

[0053] Proper control and regulation of the gas flows through the gas panel 130 is performed by mass flow controllers (not shown) and a controller unit 110 such as a computer. The showerhead 120 allows process gases from the gas panel 30 to be uniformly distributed and introduced into the process chamber 100. Illustratively, the control unit 110 comprises a central processing unit (CPU) 112, support circuitry 114, and memories containing associated control software 116. This control unit 110 is responsible for automated control of the numerous steps required for wafer processing - such as wafer transport, gas flow control, temperature control, chamber evacuation, and so on. Bi-directional communications between the control unit 110 and the various components of the apparatus 10 are handled through numerous signal cables collectively referred to as signal buses 118, some of which are illustrated in FIG. 1.

[0054] The heated pedestal 150 used in the present invention is made of aluminum, and comprises a heating element 170 embedded at a distance below the wafer support surface 151 of the pedestal 150. The heating element 170 can be made of a nickel-chromium wire encapsulated in an Incoloy sheath tube. By properly adjusting the current supplied to the heating element 170, the wafer 190 and the pedestal 150 can be maintained at a relatively constant temperature during film deposition. This is accomplished by a feedback control loop, in which the temperature of the pedestal 150 is continuously monitored by a thermocouple 172 embedded in the pedestal 150. This information is transmitted to the control unit 110 via a signal bus 118, which responds by sending the necessary signals to the heater power supply. Adjustment is subsequently made in the current supply 106 so as to maintain and control the pedestal 150 at a desirable temperature - i.e., a temperature that is appropriate for the specific process application. When the process gas mixture exits the showerhead 120, plasma enhanced thermal decomposition of the hydrocarbon compound occurs at the surface 191 of the heated wafer 190, resulting in a deposition of an amorphous carbon layer on the wafer 190.

Amorphous Carbon Layer Formation

[0055] In one embodiment of the invention the amorphous carbon layer is formed from a gas mixture of a hydrocarbon compound and an inert gas such as argon (Ar) or helium (He). The hydrocarbon compound has a general formula C_xH_y , where x has a range of between 2 and 4 and y has a range of between 2 and 10. For example, propylene (C_3H_6), propyne (C_3H_4), propane (C_3H_8), butane (C_4H_{10}), butylene (C_4H_8), butadiene (C_4H_6), or acetylene (C_2H_2) as well as combinations thereof, may be used as the hydrocarbon compound. Similarly, a variety of gases such as hydrogen (H_2), nitrogen (N_2), ammonia (NH_3), or combinations thereof, among others, may be added to the gas mixture, if desired. Ar, He, and N_2 are used to control the density and deposition rate of the amorphous carbon layer. The addition of H_2 and/or NH_3 can be used to control the hydrogen ratio of the amorphous carbon layer, as discussed below.

[0056] In general, the following deposition process parameters can be used to form the amorphous carbon layer. The process parameters range from a wafer temperature of about 100°C to about 500°C, a chamber pressure of about 1 torr to about 20 torr, a hydrocarbon gas (C_xH_y) flow rate of about 50 sccm to about 500 sccm (per 8 inch wafer), a RF power of between about 3 W/in² to about 20 W/in², and a plate spacing of between about 300 mils to about 600 mils. The above process parameters provide a typical deposition rate for the amorphous carbon layer in the range of about 100 Å/min to about 1000 Å/min and can be implemented on a 200 mm substrate in a deposition chamber available from Applied Materials, Inc.

[0057] Other deposition chambers are within the scope of the invention and the parameters listed above may vary according to the particular deposition chamber used to form the amorphous carbon layer. For example, other deposition chambers may have a larger or smaller volume, requiring gas flow rates that are larger or smaller than those recited for deposition chambers available from Applied Materials, Inc.

[0058] The as-deposited amorphous carbon layer has an adjustable carbon:hydrogen ratio that ranges from about 10 % hydrogen to about 60 % hydrogen. Controlling the hydrogen ratio of the amorphous carbon layer is desirable for tuning its optical properties as well as its etch selectivity. Specifically, as the hydrogen ratio decreases the optical properties of the as-deposited layer such as for example, the index of refraction (n) and the absorption coefficient (k) increase. Similarly, as the hydrogen ratio decreases the etch resistance of the amorphous carbon layer increases.

[0059] The light absorption coefficient, k, of the amorphous carbon layer can be varied between about 0.1 to about 1.0 at wavelengths below about 250 nm, making it suitable for use as an anti-reflective coating (ARC) at DUV wavelengths. The absorption coefficient of the

amorphous carbon layer can be varied as a function of the deposition temperature. In particular, as the temperature increases the absorption coefficient of the as-deposited layer likewise increases. For example, when propylene is the hydrocarbon compound the k value for the as-deposited amorphous carbon layers can be increased from about 0.2 to about 0.7 by increasing the deposition temperature from about 150 °C to about 480 °C.

[0060] The absorption coefficient of the amorphous carbon layer can also be varied as a function of the additive used in the gas mixture. In particular, the presence of H_2 , NH_3 , N_2 or combinations thereof, in the gas mixture can increase the k value by about 10 % to about 100 %.

Integrated Circuit Fabrication Processes

A. Amorphous Carbon Hardmask

[0061] FIGS. 2a-e illustrate schematic cross-sectional views of a substrate 200 at different stages of an integrated circuit fabrication sequence incorporating an amorphous carbon layer as a hardmask. In general, the substrate 200 refers to any workpiece on which processing is performed, and a substrate structure 250 is used to generally denote the substrate 200 together with other material layers formed on the substrate 200. Depending on the specific stage of processing, the substrate 200 may correspond to a silicon substrate, or other material layer that has been formed on the substrate. FIG. 2a, for example, illustrates a cross-sectional view of a substrate structure 250, having a material layer 202 that has been conventionally formed thereon. The material layer 202 may be an oxide (e.g., SiO_2). In general, the substrate 200 may include a layer of silicon, silicides, metals or other materials. FIG. 2a illustrates one embodiment in which the substrate 200 is silicon having a silicon dioxide layer formed thereon.

[0062] FIG. 2b depicts an amorphous carbon layer 204 deposited on the substrate structure 250 of FIG. 2a. The amorphous carbon layer 204 is formed on the substrate structure 250 according to the process parameters described above. The thickness of the amorphous carbon layer is variable depending on the specific stage of processing. Typically, the amorphous carbon layer has a thickness in the range of about 50 Å to about 1000 Å.

[0063] Dependant on the etch chemistry of the energy sensitive resist material used in the fabrication sequence, an intermediate layer 206 is formed on the amorphous carbon layer 204. The intermediate layer 206 functions as a mask for the amorphous carbon layer 204 when the pattern is transferred therein. The intermediate layer 206 is conventionally formed on the amorphous carbon layer 204. The intermediate layer 206 may be an oxide, nitride, silicon oxynitride, silicon carbide, amorphous silicon or other materials.

[0064] A layer of energy sensitive resist material 208

is formed on the intermediate layer 206. The layer of energy sensitive resist material 208 can be spin coated on the substrate to a thickness within the range of about 2000 Å to about 6000 Å. Most energy sensitive resist materials are sensitive to ultraviolet (UV) radiation having a wavelength less than about 450 nm. DUV resist materials are sensitive to UV radiation having wavelengths of 245 nm or 193 nm.

[0065] An image of a pattern is introduced into the layer of energy sensitive resist material 208 by exposing such energy sensitive resist material 208 to UV radiation via mask 210. The image of the pattern introduced in the layer of energy sensitive resist material 208, is developed in an appropriate developer to define the pattern through such layer as shown in FIG. 2c. Thereafter, referring to FIG. 2d, the pattern defined in the energy sensitive resist material 208 is transferred through both the intermediate layer 206 and the amorphous carbon layer 204. The pattern is transferred through the intermediate layer 206 using the energy sensitive resist material 208 as a mask. The pattern is transferred through the intermediate layer 206 by etching the intermediate layer 206 using an appropriate chemical etchant. The pattern is then transferred through the amorphous carbon layer 204 using the intermediate layer 206 as a mask. The pattern is transferred through the amorphous carbon layer 204 by etching the amorphous carbon layer 204 using an appropriate chemical etchant (e. g., ozone, oxygen or ammonia plasmas).

[0066] FIG. 2e illustrates the completion of the integrated circuit fabrication sequence by the transfer of the pattern defined in the amorphous carbon layer 204 through the silicon dioxide layer 202 using the amorphous carbon layer 204 as a hardmask.

[0067] After the silicon dioxide layer 202 is patterned, the amorphous carbon layer 204 can optionally be stripped from the substrate 200 by etching it in an ozone, oxygen or ammonia plasma.

[0068] In a specific example of a fabrication sequence, the pattern defined in the amorphous carbon hardmask can be incorporated into the structure of the integrated circuit, such as a damascene structure. Damascene structures are typically used to form metal interconnects on integrated circuits.

[0069] FIGS. 3a-3e illustrate schematic cross-sectional views of a substrate 260 at different stages of a damascene structure fabrication sequence incorporating an amorphous carbon layer therein. Depending on the specific stage of processing, substrate 260 may correspond to a silicon substrate, or other material layer that has been formed on the substrate. FIG. 3a, for example, illustrates a cross-sectional view of a substrate 260 having a dielectric layer 262 formed thereon. The dielectric layer 262 may be an oxide (e. g., silicon dioxide, fluorosilicate glass). In general, the substrate 260 may include a layer of silicon, silicides, metals or other materials.

[0070] FIG. 3a illustrates one embodiment in which

the substrate 260 is silicon having a fluorosilicate glass layer formed thereon. The dielectric layer 262 has a thickness of about 5,000 Å to about 10,000 Å, depending on the size of the structure to be fabricated. An amorphous carbon layer 264 is formed on the dielectric layer 262. The amorphous carbon layer is formed on the dielectric layer 262 according to the process parameters described above. The amorphous carbon layer 264 has a thickness of about 200 Å to about 1000 Å.

[0071] Referring to FIG. 3b, the amorphous carbon layer 264 is patterned and etched to define contact/via openings 266 and to expose the dielectric layer 262, in areas where the contacts/vias are to be formed. The amorphous carbon layer 264 is patterned using conventional lithography and etched using oxygen or ammonia plasmas.

[0072] The contact/via openings 266 formed in the amorphous carbon layer 264 are then transferred into the dielectric layer 262 using the amorphous carbon layer 264 as a hard mask as shown in FIG. 3c. The contacts/vias 266 are etched using reactive ion etching or other anisotropic etching techniques. After the contacts/vias 266 are transferred into the dielectric layer 262, the amorphous carbon layer is stripped from dielectric layer 262 by etching it in an ozone, oxygen or ammonia plasma as depicted in FIG. 3d.

[0073] Referring to FIG. 3e, a metallization structure is formed in the contacts/vias 266 using a conductive material 274 such as aluminum, copper, tungsten, or combinations thereof. Typically, copper is used to form the metallization structure due to its low resistivity (about 1.7 $\mu\Omega\text{-cm}$). The conductive material 274 is deposited using chemical vapor deposition, physical vapor deposition, electroplating, or combinations thereof, to form the damascene structure. Preferably, a barrier layer 272 such as tantalum, tantalum nitride, or other suitable barrier is first deposited conformally in the metallization structure to prevent metal migration into the surrounding dielectric material layer 262. Additionally, the dielectric layer 262 preferably has a low dielectric constant (dielectric constants less than about 4.5) so as to prevent capacitive coupling between adjacent contacts/vias 266 of the metallization structure.

B. Amorphous Carbon Anti-Reflective Coating (ARC)

[0074] FIGS. 4a-4c illustrate schematic cross-sectional views of a substrate 300 at different stages of an integrated circuit fabrication sequence incorporating an amorphous carbon layer as an anti-reflective coating (ARC). In general, the substrate 300 refers to any workpiece on which film processing is performed, and a substrate structure 350 is used to generally denote the substrate 300 together with other material layers formed on the substrate 300. Depending on the specific stage of processing, substrate 300 may correspond to a silicon substrate, or other material layer, which has been formed on the substrate. FIG. 4a, for example, illustrates

a cross-sectional view of a substrate structure 350 in which the substrate 300 is an oxide layer formed on a silicon wafer.

[0075] An amorphous carbon layer 302 is formed on the substrate 300. The amorphous carbon layer 302 is formed on the substrate 300 according to the process parameters described above. The amorphous carbon layer has a refractive index (n) in the range of about 1.5 to 1.9 and an absorption coefficient (k) in the range of about 0.1 to about 1.0 at wavelengths less than about 250 nm making it suitable for use as an ARC at DUV wavelengths. The refractive index (n) and absorption coefficient (k) for the amorphous carbon ARC are tunable, in that they can be varied in the desired range as a function of the temperature as well as the composition of the gas mixture during layer formation. The thickness of the amorphous carbon layer is variable depending on the specific stage of processing. Typically, the amorphous carbon layer has a thickness of about 200 Å to about 1100 Å.

[0076] FIG. 4b depicts a layer of energy sensitive resist material 304 formed on the substrate structure 350 of FIG. 4a. The layer of energy sensitive resist material can be spin coated on the substrate to a thickness within the range of about 2000 Å to about 6000 Å. The energy sensitive resist material is sensitive to DUV radiation having a wavelength less than 250 nm.

[0077] An image of a pattern is introduced into the layer of energy sensitive resist material 304 by exposing such energy sensitive resist material 304 to DUV radiation via mask 306. The image of the pattern introduced into the layer of energy sensitive resist material 304 is developed in an appropriate developer to define the pattern through such layer. Thereafter, referring to FIG. 4c, the pattern defined in the energy sensitive resist material 304 is transferred through the amorphous carbon layer 302. The pattern is transferred through the amorphous carbon layer 302 using the energy sensitive resist material 304 as a mask. The pattern is transferred through the amorphous carbon layer 302 by etching it using an appropriate chemical etchant (e. g., ozone, oxygen or ammonia plasmas).

[0078] After the amorphous carbon 302 is patterned, such pattern is optionally transferred into the substrate 300. Typically, when substrate 300 comprises an oxide layer on a silicon substrate, the etch selectivity of the oxide to a resist mask is about 3:1 to about 5:1. Specifically, the oxide will etch about 3 to 5 times faster than the resist. In contrast, the amorphous carbon ARC layer of the present invention has an etch selectivity to the oxide of greater than about 10:1. That is, the oxide will etch more than 10 times faster than the amorphous carbon ARC. Thus, the amorphous carbon ARC layer also provides greater etch selectivity as a hardmask for patterning the oxide, without the added complexity of requiring an additional intermediate hardmask layer.

[0079] In an alternate embodiment, the amorphous carbon layer can have an absorption coefficient (k) that

varies across the thickness of the layer. That is, the amorphous carbon layer can have an absorption coefficient gradient formed therein. Such a gradient is formed as a function of the temperature and the composition of the gas mixture during layer formation.

[0080] At any interface between two material layers, reflections can occur because of differences in their refractive indices (n) and absorption coefficients (k). When the amorphous carbon ARC has a gradient, it is possible to match the refractive indices (n) and the absorption coefficients (k) of the two material layers so there is minimal reflection and maximum transmission into the amorphous carbon ARC. Then the refractive index (n) and absorption coefficient (k) of the amorphous carbon ARC can be gradually adjusted to absorb all of the light transmitted therein.

C. Multi-Layer Amorphous Carbon Anti-Reflective Coating (ARC)

[0081] FIGS. 5a-5d illustrate schematic cross-sectional views of a substrate 400 at different stages of an integrated circuit fabrication sequence incorporating a multi-layer amorphous carbon anti-reflective coating (ARC) structure. In general, the substrate 400 refers to any workpiece on which film processing is performed, and a substrate structure 450 is used to generally denote the substrate 400 together with other material layers formed on the substrate 400. Depending on the specific stage of processing, substrate 400 may correspond to a silicon substrate, or other material layer, which has been formed on the substrate. FIG. 5a, for example, illustrates a cross-sectional view of a substrate structure 450 in which the substrate 400 is a silicon wafer.

[0082] A first amorphous carbon layer 402 is formed on the substrate 400. The first amorphous carbon layer 402 is formed on the substrate 400 according to the process parameters described above. The first amorphous carbon layer 402 is designed primarily for light absorption. As such, the first amorphous carbon layer 402 has an index of refraction in the range of about 1.5 to about 1.9 and an absorption coefficient (k) in the range of about 0.5 to about 1.0 at wavelengths less than about 250 nm. The thickness of the first amorphous carbon layer 402 is variable depending on the specific stage of processing. Typically, the first amorphous carbon layer 402 has a thickness in the range of about 300 Å to about 1500 Å.

[0083] A second amorphous carbon layer 404 is formed on the first amorphous carbon layer 402. The second amorphous carbon layer 404 is also formed according to the process parameters described above. The second amorphous carbon layer 404 is designed primarily for phase shift cancellation. Specifically, the second amorphous carbon layer is designed to create reflections that cancel those generated at the interface with an overlying material layer (e. g., an energy sensitive resist material). As such, the second amorphous

carbon layer 404 has an index of refraction of about 1.5 to about 1.9 and an absorption coefficient in the range of about 0.1 to about 0.5.

[0084] The thickness of the second amorphous carbon layer 404 is also variable depending on the specific stage of processing. Typically, the second amorphous carbon layer 404 has a thickness in the range of about 300 Å to about 700 Å. The refractive index (n) and the absorption coefficient (k) of the first and second amorphous carbon layers are tunable, in that they can be varied as a function of the temperature as well as the composition of the gas mixture during layer formation.

[0085] Additional amorphous carbon layers may be included in the multi-layered amorphous carbon ARC structure. For example, more than one top layer can be used to cancel reflections generated at the interface with for example, the energy sensitive resist material, while more than one bottom layer can be used to absorb light transmitted therein minimizing reflections at the interface of the multi-layered amorphous carbon ARC structure with underlying materials such as for example low dielectric constant oxides.

[0086] FIG. 5b depicts a layer of energy sensitive resist material 406 formed on the substrate structure 450 of FIG. 5a. The layer of energy sensitive resist material can be spin coated on the substrate to a thickness within the range of about 2000 Å to about 6000 Å. The energy sensitive resist material is sensitive to DUV radiation having a wavelength less than 250 nm.

[0087] An image of a pattern is introduced into the layer of energy sensitive resist material 406 by exposing such energy sensitive resist material 406 to DUV radiation via mask 408.

[0088] The image of the pattern introduced into the layer of energy sensitive resist material 406 is developed in an appropriate developer to define the pattern through such layer as shown in FIG. 5c. Thereafter, referring to FIG. 5d, the pattern defined in the energy sensitive resist material 406 is transferred through both amorphous carbon layers 404, 402 using the energy sensitive resist material 406 as a mask. The pattern is transferred through the amorphous carbon layers 404, 402 by etching them using an appropriate chemical etchant (e. g., ozone, oxygen or ammonia plasmas). After the multi-layer ARC is patterned, such pattern is optionally transferred into the substrate.

[0089] The multi-layered amorphous carbon ARC structure described with reference to FIGS. 5a-5d, also provides etch selectivity as a hardmask for patterning such underlying material layers as low dielectric constant oxides, without the added complexity of requiring an additional intermediate hardmask layer, as discussed previously for the single-layer amorphous carbon ARC.

Claims

1. A method of forming an amorphous carbon layer on a substrate comprising:

positioning a substrate in a deposition chamber;
providing a gas mixture to the deposition chamber, wherein the gas mixture comprises one or more hydrocarbon compounds and an inert gas; and
heating the gas mixture to thermally decompose the one or more hydrocarbon compounds in the gas mixture to form an amorphous carbon layer on the substrate.

2. A method of forming a device, comprising:

forming one or more amorphous carbon layers on a substrate; and
defining a pattern in at least one region of the one or more amorphous carbon layers.

3. The method of claim 2 further comprising transferring the pattern defined in the at least one region of the one or more amorphous carbon layers into the substrate using the one or more amorphous carbon layers as a mask.

4. The method of claim 2 wherein definition of the pattern in the at least one region of the one or more amorphous carbon layers, comprises:

forming a layer of energy sensitive resist material on the intermediate layer;
introducing an image of the pattern into the layer of energy sensitive resist material by exposing the energy sensitive resist material to patterned radiation;
developing the image of the pattern introduced into the layer of energy sensitive resist material; and
transferring the pattern through the one or more amorphous carbon layers using the layer of energy sensitive resist material as a mask.

5. The method of claim 4 further comprising:

forming an intermediate layer on the one or more amorphous carbon layers prior to forming the layer of energy sensitive resist thereon, introducing the image of a pattern therein, and developing the image of the pattern;
transferring the image of the pattern developed in the layer of energy sensitive resist material through the intermediate layer using the energy sensitive resist material as a mask; and
transferring the pattern through the one or more

amorphous carbon layers using the intermediate layer as a mask.

6. The method of claim 5 wherein the intermediate layer is an oxide.

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7. The method of claim 2 wherein the one or more amorphous carbon layers are anti-reflective coatings at wavelengths less than 250 nm.

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8. The method of claim 40 wherein the one or more amorphous carbon layers each have an absorption coefficient in the range of 0.1 to 1.0 at wavelengths less than 250 nm.

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9. The method of claim 8 wherein the absorption coefficient varies across the thickness of the one or more amorphous carbon layers from 0.1 to 1.0.

10. A method of fabricating a damascene structure, comprising

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forming a dielectric layer on a substrate;
forming an amorphous carbon layer on the dielectric layer;
patterning the amorphous carbon layer to define contacts/vias therethrough;
transferring the pattern formed in the amorphous carbon layer through the dielectric layer to form contacts/vias therein;
removing the amorphous carbon layer from the patterned dielectric layer; and
filling the contacts/vias formed in the dielectric layer with a conductive material.

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11. The method of claim 10 wherein the amorphous carbon layer is formed on the dielectric layer by

positioning the substrate in a deposition chamber, wherein the substrate has a dielectric layer thereon;
providing a gas mixture to the deposition chamber, wherein the gas mixture comprises one or more hydrocarbon compounds and an inert gas; and
heating the gas mixture to thermally decompose the one or more hydrocarbon compounds in the gas mixture to form the amorphous carbon layer on the dielectric layer.

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12. The method of any of the preceding claims wherein the one or more hydrocarbon compounds in the gas mixture have the general formula C_xH_y , wherein x has a range of 2 to 4 and y has a range of 2 to 10.

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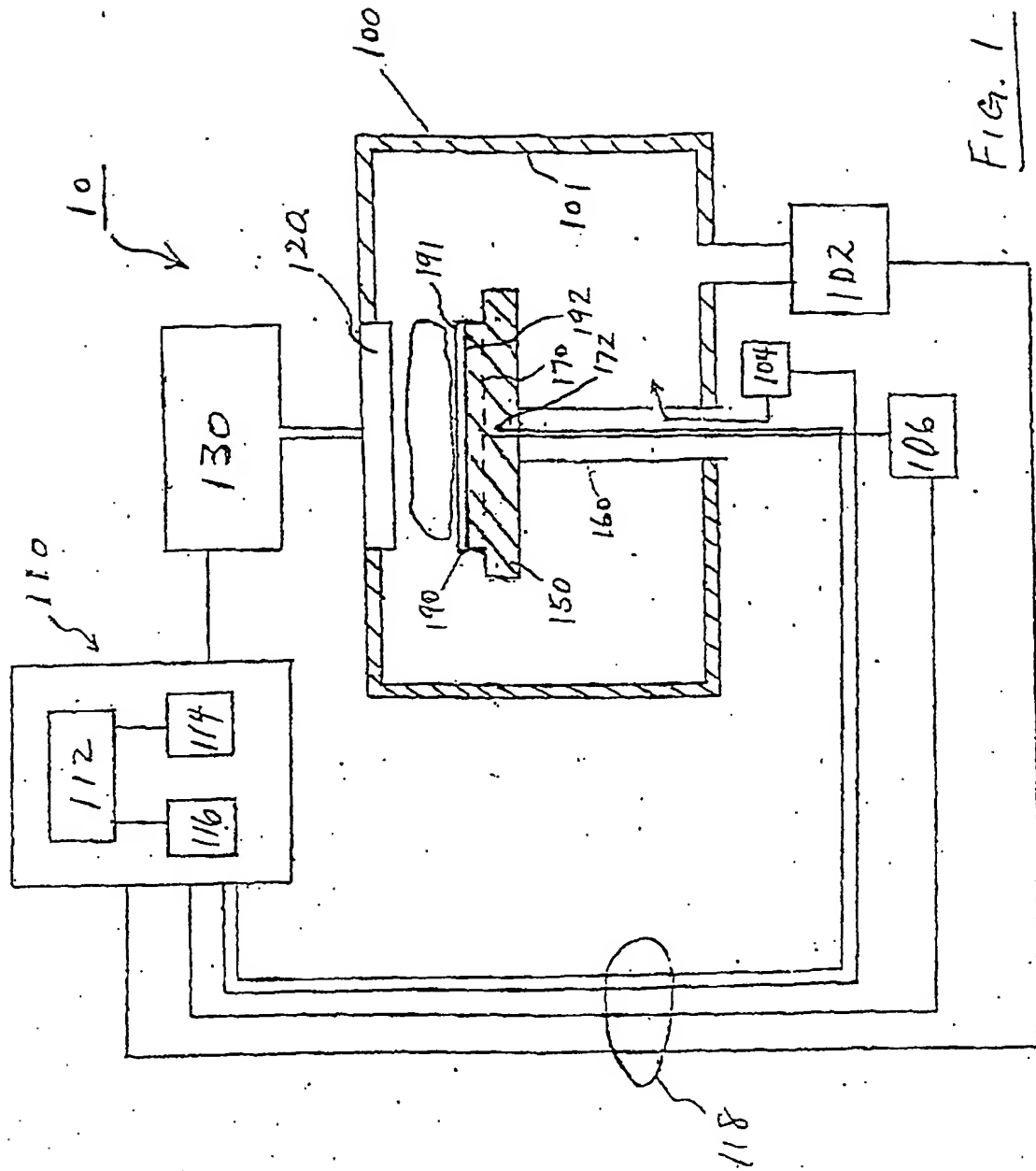
13. The method of any of the preceding claims wherein the amorphous carbon layer has a carbon:hydrogen ratio in the range of 10 % hydrogen to 60 %

hydrogen.

14. The method of any of the preceding claims wherein the gas mixture is heated by applying an electric field thereto.

15. The method of any of the preceding claims wherein the electric field is a RF field the the RF power of which is in a range of 3 W/in² to 20 W/in².

16. A computer storage medium containing a software routine that, when executed, causes a general purpose computer to control a deposition chamber using a method of any of the preceding claims.



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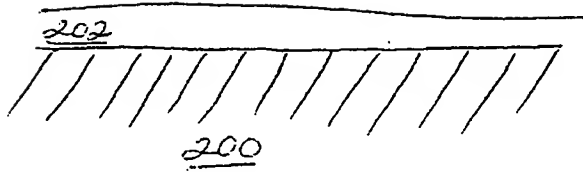


Fig. 2a

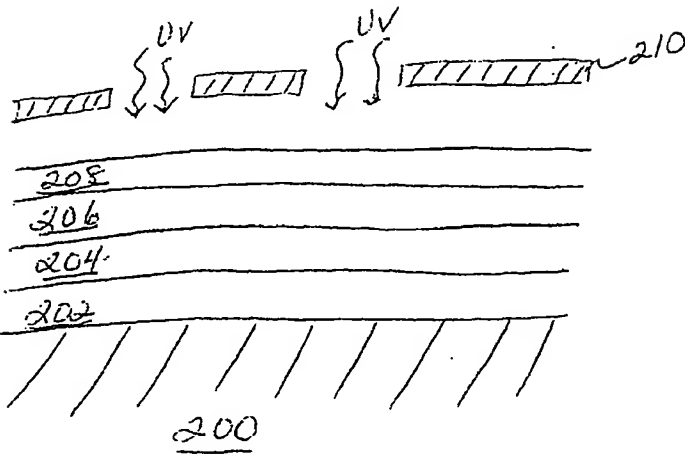


Fig. 2b

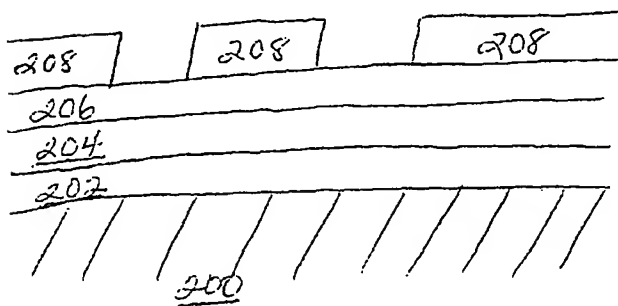


Fig. 2c

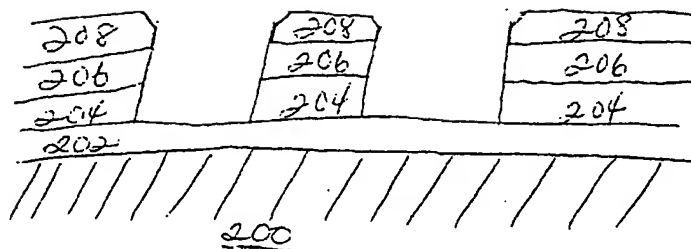


Fig. 2d

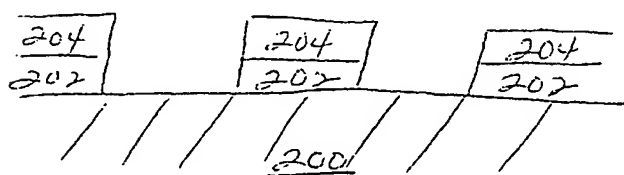


Fig. 2e

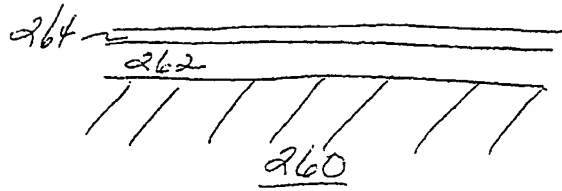


Fig. 3a

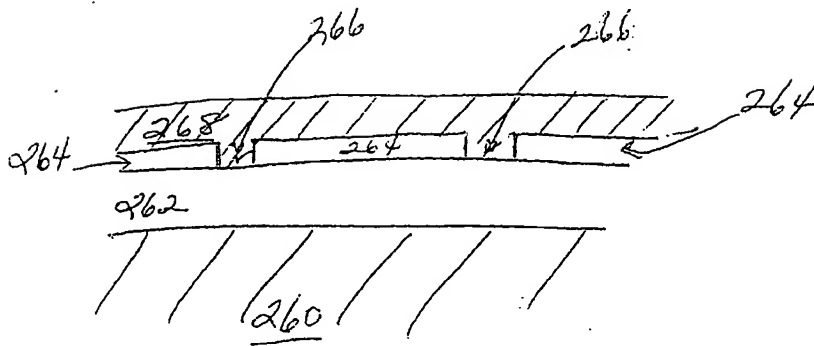


Fig. 3b

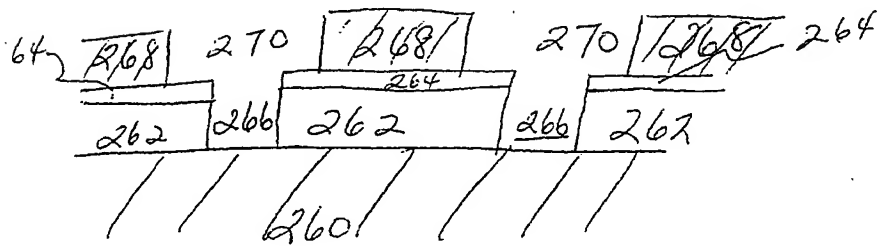


Fig. 3c

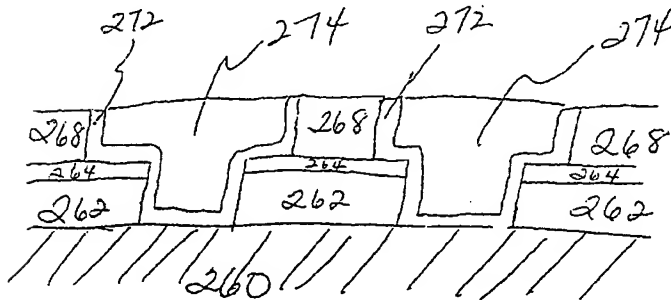


Fig. 3d

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Fig. 4a

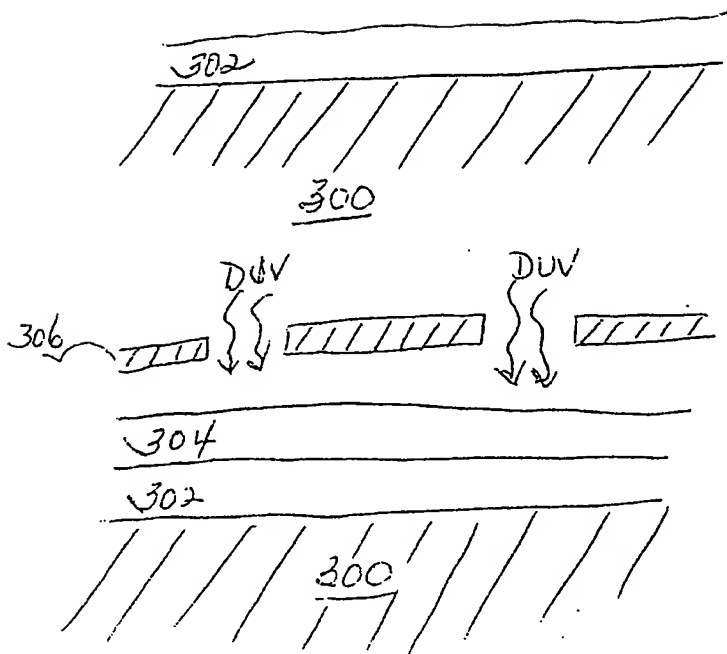


Fig. 4b

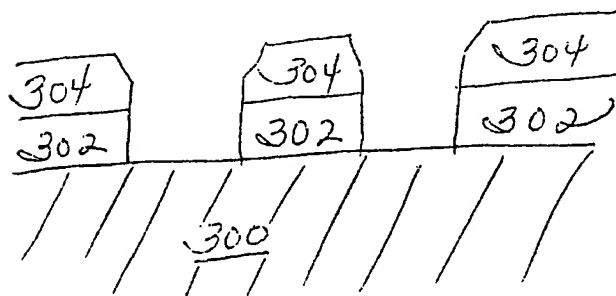


Fig. 4c

450

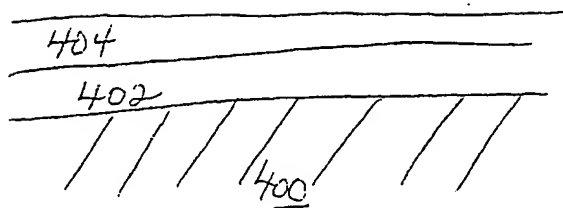


Fig. 5a

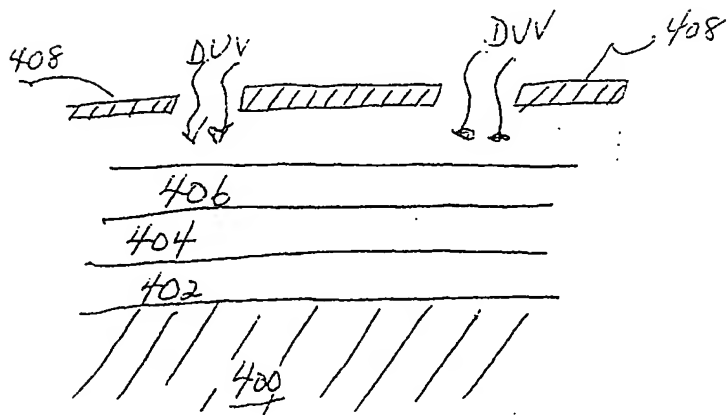


Fig. 5b

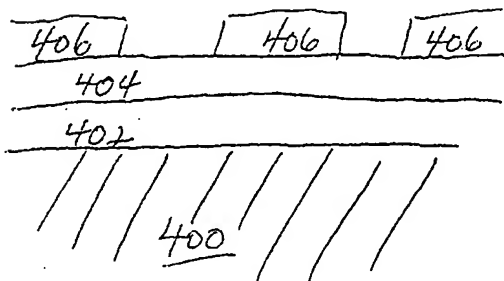


Fig. 5c

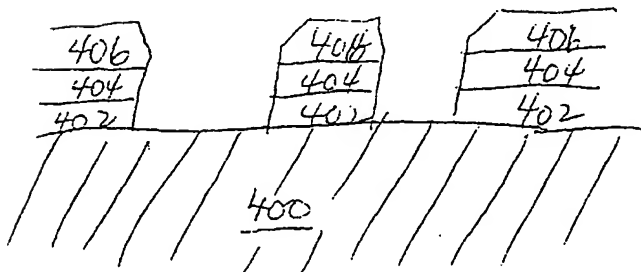


Fig. 5d